

05/14/2006 10669301a.trn

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Take survey: <http://www.zoomerang.com/survey.zgi?p=WEB2259HNKWTUW>

Thank you in advance for your participation.

FILE 'HOME' ENTERED AT 09:58:49 ON 14 MAY 2006

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THIS COMMAND NOT AVAILABLE IN THE CURRENT FILE

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Choice (Y/n) :

Switching to the Registry File...
Some commands only work in certain files. For example, the EXPAND command can only be used to look at the index in a file which has an index. Enter "HELP COMMANDS" at an arrow prompt (=>) for a list of commands which can be used in this file.

=> FILE REGISTRY

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

FILE 'REGISTRY' ENTERED AT 09:59:00 ON 14 MAY 2006
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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 12 MAY 2006 HIGHEST RN 884047-29-4
DICTIONARY FILE UPDATES: 12 MAY 2006 HIGHEST RN 884047-29-4

New CAS Information Use Policies. enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH January 6, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

* * *

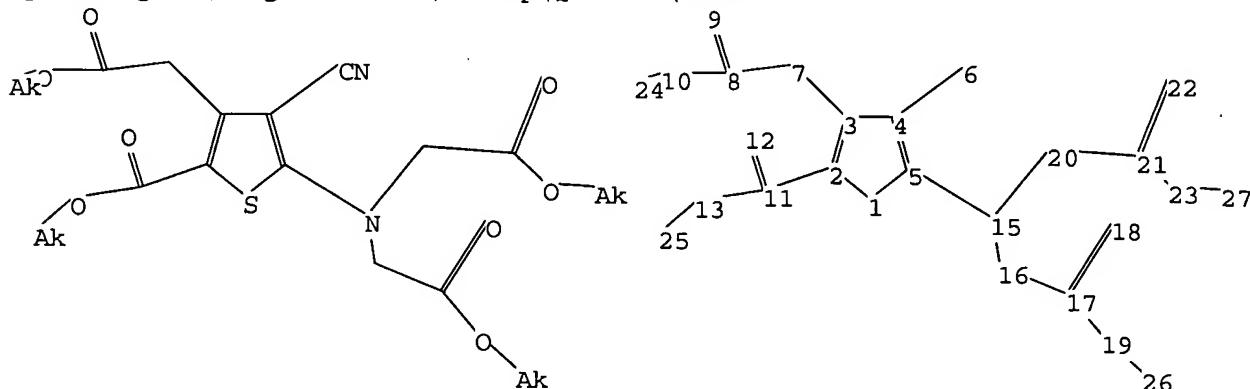
* The CA roles and document type information have been removed from *
 * the IDE default display format and the ED field has been added, *
 * effective March 20, 2005. A new display format, IDERL, is now *
 * available and contains the CA role and document type information. *
 *

Structure search iteration limits have been increased. See HELP SLIMITS for details.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

=>
 Uploading C:\Program Files\Stnexp\Queries\10669301a.str



chain nodes :
 6 7 8 9 10 11 12 13 15 16 17 18 19 20 21 22 23 24 25 26 27
 ring nodes :
 1 2 3 4 5
 chain bonds :
 2-11 3-7 4-6 5-15 7-8 8-9 8-10 10-24 11-12 11-13 13-25 15-16 15-20
 16-17 17-18 17-19 19-26 20-21 21-22 21-23 23-27
 ring bonds :
 1-2 1-5 2-3 3-4 4-5
 exact/norm bonds :
 5-15 8-9 8-10 10-24 11-12 11-13 13-25 15-16 15-20 17-18 17-19 19-26
 21-22 21-23 23-27
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 1-2 1-5 2-3 2-11 3-4 3-7 4-5 4-6 7-8 16-17 20-21
 isolated ring systems :
 containing 1 :

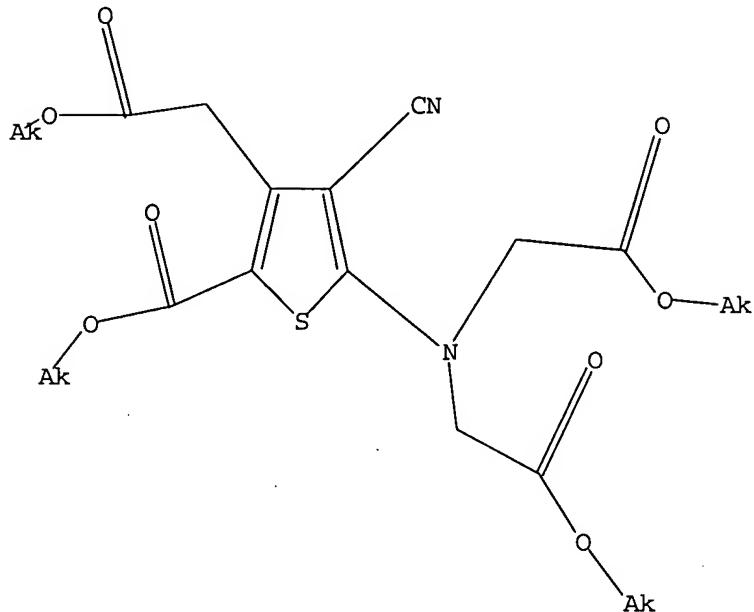
Match level :

05/14/2006 10669301a.trn

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:CLASS 7:CLASS 8:CLASS 9:CLASS
10:CLASS 11:CLASS 12:CLASS 13:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS
19:CLASS 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS 26:CLASS
27:CLASS

L1 STRUCTURE UPLOADED

=> d 11
L1 HAS NO ANSWERS
L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s 11
SAMPLE SEARCH INITIATED 09:59:16 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 2 TO ITERATE

100.0% PROCESSED 2 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 2 TO 124
PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s 11 sss full
FULL SEARCH INITIATED 09:59:22 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 55 TO ITERATE

100.0% PROCESSED 55 ITERATIONS

3 ANSWERS

05/14/2006 10669301a.trn

SEARCH TIME: 00.00.01

L3 3 SEA SSS FUL L1

=> s 13 and strontium hydroxide
67645 STRONTIUM
37379 HYDROXIDE
85 HYDROXIDES
37379 HYDROXIDE
(HYDROXIDE OR HYDROXIDES)
331 STRONTIUM HYDROXIDE
(STRONTIUM(W) HYDROXIDE)

L4 0 L3 AND STRONTIUM HYDROXIDE

=> FIL HCAPLUS

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	176.90	177.11

FILE 'HCAPLUS' ENTERED AT 10:00:06 ON 14 MAY 2006
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 14 May 2006 VOL 144 ISS 21
FILE LAST UPDATED: 12 May 2006 (20060512/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d his

(FILE 'HOME' ENTERED AT 09:58:49 ON 14 MAY 2006)

FILE 'REGISTRY' ENTERED AT 09:59:00 ON 14 MAY 2006

L1 STRUCTURE UPLOADED
L2 0 S L1
L3 3 S L1 SSS FULL
L4 0 S L3 AND STRONTIUM HYDROXIDE

FILE 'HCAPLUS' ENTERED AT 10:00:06 ON 14 MAY 2006

=> s 13
L5 4 L3
=> s 15 and strontium hydroxide

184117 STRONTIUM
 4 STRONTIUMS
 184118 STRONTIUM
 (STRONTIUM OR STRONTIUMS)
 274546 HYDROXIDE
 45454 HYDROXIDES
 296569 HYDROXIDE
 (HYDROXIDE OR HYDROXIDES)
 1296 STRONTIUM HYDROXIDE
 (STRONTIUM(W) HYDROXIDE)
 1 L5 AND STRONTIUM HYDROXIDE

L6

=> d 16 ibib abs hitstr tot

L6 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2004:249307 HCAPLUS
 DOCUMENT NUMBER: 140:272696
 TITLE: New process for industrial synthesis of strontium
 ranelate and its hydrates
 INVENTOR(S): Vaysse, Ludot Lucile; Lecouve, Jean Pierre; Langlois,
 Pascal
 PATENT ASSIGNEE(S): Les Laboratoires Servier, Fr.
 SOURCE: Fr. Demande, 22 pp.
 CODEN: FRXXBL
 DOCUMENT TYPE: Patent
 LANGUAGE: French
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
FR 2844795	A1	20040326	FR 2002-11763	20020924
FR 2844795	B1	20041022		
EP 1403266	A1	20040331	EP 2003-292319	20030922
R: AT, BE, CH, DE, DK, ES, FR, IE, SI, LT, LV, FI, RO, MK		GB, GR, IT, LI, LU, NL, SE, MC, PT, CY, AL, TR, BG, CZ, EE, HU, SK		
AU 2003248281	A1	20040408	AU 2003-248281	20030922
WO 2004029036	A1	20040408	WO 2003-FR2777	20030922
W: AE, AG, AL, AM, AT, AU, AZ, CO, CR, CU, CZ, DE, DK, DM, GM, HR, HU, ID, IL, IN, IS, LS, LT, LU, LV, MA, MD, MG, PG, PH, PL, PT, RO, RU, SC, TR, TT, TZ, UA, UG, US, UZ, RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG		BA, BB, BG, BR, BY, BZ, CA, CH, CN, DZ, EC, EE, ES, FI, GB, GD, GE, GH, KP, KR, KZ, LC, LK, LR, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, ZA, ZM, ZW		
AU 2003282179	A1	20040419	AU 2003-282179	20030922
JP 2004149516	A2	20040527	JP 2003-330440	20030922
CA 2442878	AA	20040324	CA 2003-2442878	20030923
NO 2003004235	A	20040325	NO 2003-4235	20030923
ZA 2003007409	A	20040707	ZA 2003-7409	20030923
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BR 2003004213	A	20040831	BR 2003-4213	20030923
US 2004063972	A1	20040401	US 2003-669301	20030924
CN 1496986	A	20040519	CN 2003-134813	20030924
SG 110071	A1	20050428	SG 2003-5555	20030924
HK 1065791	A1	20051014	HK 2004-108552	20041101

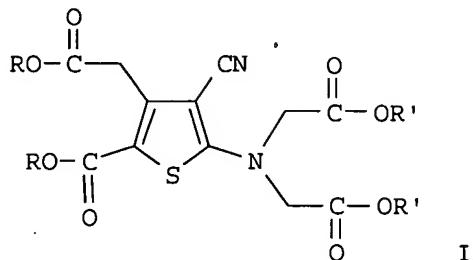
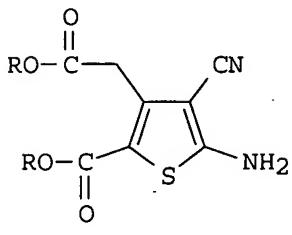
PRIORITY APPLN. INFO.:

FR 2002-11763
WO 2003-FR2777A 20020924
W 20030922

OTHER SOURCE(S):

MARPAT 140:272696

GI



AB An industrial process for the synthesis of strontium ranelate and its hydrates consists of: reaction of RO₂CCH₂COCH₂CO₂R (R = linear or branched C₁-6 alkyl) with malononitrile (NCCH₂CN) in MeOH in presence of morpholine (>0.95 mol per mol diester) to give the morpholinium salt of ROCOCH₂C[:C(CN)2]CH:CO₂R-, followed by refluxing with sulfur to give thiophene derivative I (same R). Reaction of the latter (as diacid) with BrCH₂CO₂R' (R' = e.g., Me or Et) in the presence of a catalytic quantity of C₈-10 quaternary ammonium salt and K₂CO₃ in an organic solvent at reflux affords tetracarboxylate II, which reacts with Sr(OH)₂ at reflux in water for ≥ 5 h to give strontium ranelate and its hydrates. Thus, the octahydrate of strontium ranelate was prepared by this method (96% yield and 98% purity in final step).

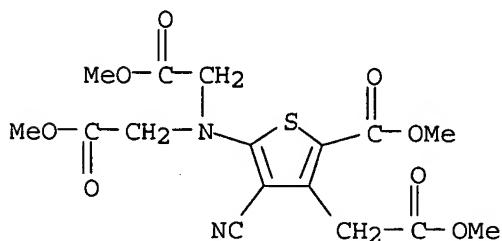
IT 674773-13-8P

RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)

(for industrial preparation of strontium ranelate and its hydrates)

RN 674773-13-8 HCAPLUS

CN 3-Thiopheneacetic acid, 5-[bis(2-methoxy-2-oxoethyl)amino]-4-cyano-2-(methoxycarbonyl)-, methyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT:

3

THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d 15 ibib abs hitstr tot

L5 ANSWER 1 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN

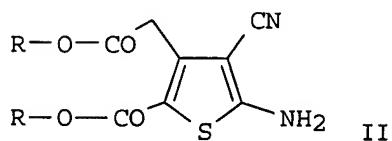
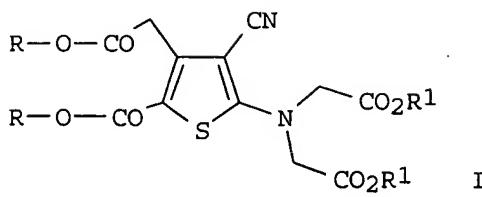
ACCESSION NUMBER: 2004:252227 HCAPLUS

DOCUMENT NUMBER: 140:270729

TITLE: Process for the industrial synthesis of tetraesters of 5-[bis(carboxymethyl)amino]-3-carboxymethyl-4-cyano-2-thiophenecarboxylic acid and their application to the synthesis of bivalent salts of ranelic acid and their hydrates
 INVENTOR(S): Vaysse-Ludot, Lucile; Lecouve, Jean-pierre; Langlois, Rascal
 PATENT ASSIGNEE(S):
 SOURCE: U.S. Pat. Appl. Publ., 4 pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004059134	A1	20040325	US 2003-669302	20030924
FR 2844797	A1	20040326	FR 2002-11765	20020924
FR 2844797	B1	20041022		
EP 1403265	A1	20040331	EP 2003-292318	20030922
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AU 2003248280	A1	20040408	AU 2003-248280	20030922
WO 2004029034	A1	20040408	WO 2003-FR2775	20030922
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	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG		
AU 2003299095	A1	20040419	AU 2003-299095	20030922
JP 2004269496	A2	20040930	JP 2003-330439	20030922
CA 2442881	AA	20040324	CA 2003-2442881	20030923
NO 2003004236	A	20040325	NO 2003-4236	20030923
NZ 528401	A	20040528	NZ 2003-528401	20030923
ZA 2003007411	A	20040707	ZA 2003-7411	20030923
BR 2003004203	A	20040824	BR 2003-4203	20030923
CN 1500784	A	20040602	CN 2003-134812	20030924
SG 110069	A1	20050428	SG 2003-5553	20030924
PRIORITY APPLN. INFO.:			FR 2002-11765	A 20020924
			WO 2003-FR2775	W 20030922

OTHER SOURCE(S): CASREACT 140:270729; MARPAT 140:270729
 GI



AB Tetraesters of 5-[bis(carboxymethyl)amino]-3-carboxymethyl-4-cyano-2-thiophenecarboxylic acid [I; R, R1 = (un)branched C1-6 alkyl] are prepared in high yield and selectivity by the alkylation of the corresponding 5-amino compound (II) with an alkyl bromoacetate ester $\text{BrCH}_2\text{CO}_2\text{R1}$ in the presence of a catalytic amount of a quaternary ammonium compound, potassium carbonate acid scavenger at reflux in an organic solvent, the reaction mixture is then concentrated by distillation, an a nonsolvent added to cause precipitation of the product with cooling. The synthesis of bivalent salts of ranelic acid, and especially strontium ranelate and its hydrates, is claimed.

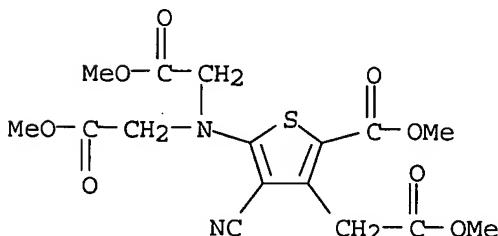
IT 674773-13-8P 674800-87-4P

RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)

(process for the industrial synthesis of tetraesters of 5-[bis(carboxymethyl)amino]-3-carboxymethyl-4-cyano-2-thiophenecarboxylic acid and their application to the synthesis of bivalent salts of ranelic acid and their hydrates)

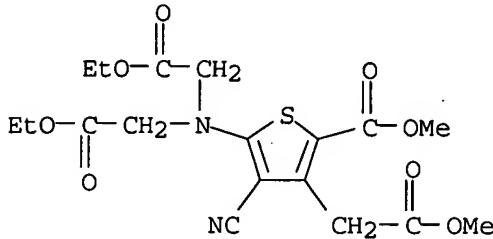
RN 674773-13-8 HCPLUS

CN 3-Thiopheneacetic acid, 5-[bis(2-methoxy-2-oxoethyl)amino]-4-cyano-2-(methoxycarbonyl)-, methyl ester (9CI) (CA INDEX NAME)



RN 674800-87-4 HCPLUS

CN 3-Thiopheneacetic acid, 5-[bis(2-ethoxy-2-oxoethyl)amino]-4-cyano-2-(methoxycarbonyl)-, methyl ester (9CI) (CA INDEX NAME)

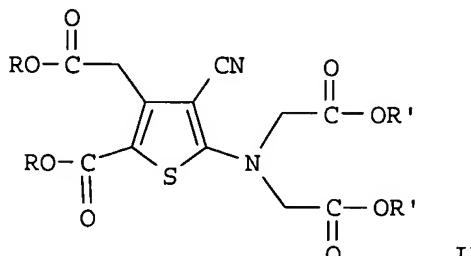
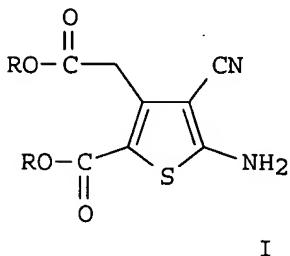


L5 ANSWER 2 OF 4 HCPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2004:249307 HCPLUS
 DOCUMENT NUMBER: 140:272696
 TITLE: New process for industrial synthesis of strontium ranelate and its hydrates
 INVENTOR(S): Vaysse, Ludot, Lucile; Lecouve, Jean Pierre; Langlois, Pascal
 PATENT ASSIGNEE(S): Les Laboratoires Servier, Fr.
 SOURCE: Fr. Demande, 22 pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: French
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
FR 2844795	A1	20040326	FR 2002-11763	20020924
FR 2844795	B1	20041022		
EP 1403266	A1	20040331	EP 2003-292319	20030922
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
AU 2003248281	A1	20040408	AU 2003-248281	20030922
WO 2004029036	A1	20040408	WO 2003-FR2777	20030922
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2003282179	A1	20040419	AU 2003-282179	20030922
JP 2004149516	A2	20040527	JP 2003-330440	20030922
CA 2442878	AA	20040324	CA 2003-2442878	20030923
NO 2003004235	A	20040325	NO 2003-4235	20030923
ZA 2003007409	A	20040707	ZA 2003-7409	20030923
NZ 528402	A	20040730	NZ 2003-528402	20030923
BR 2003004213	A	20040831	BR 2003-4213	20030923
US 2004063972	A1	20040401	US 2003-669301	20030924
CN 1496986	A	20040519	CN 2003-134813	20030924

SG 110071	A1 20050428	SG 2003-5555	20030924
HK 1065791	A1 20051014	HK 2004-108552	20041101
PRIORITY APPLN. INFO.:		FR 2002-11763	A 20020924
		WO 2003-FR2777	W 20030922

OTHER SOURCE(S): MARPAT 140:272696
GI



AB An industrial process for the synthesis of strontium ranelate and its hydrates consists of: reaction of $\text{RO}_2\text{CCH}_2\text{COCH}_2\text{CO}_2\text{R}$ (R = linear or branched C1-6 alkyl) with malononitrile (NCCH_2CN) in MeOH in presence of morpholine (>0.95 mol per mol diester) to give the morpholinium salt of $\text{ROCOCH}_2\text{C}[:\text{C}(\text{CN})_2]\text{CH}:\text{C}(\text{OR})\text{O}^-$, followed by refluxing with sulfur to give thiophene derivative I (same R). Reaction of the latter (as diacid) with $\text{BrCH}_2\text{CO}_2\text{R}'$ (R' = e.g., Me or Et) in the presence of a catalytic quantity of C8-10 quaternary ammonium salt and K_2CO_3 in an organic solvent at reflux affords tetracarboxylate II, which reacts with $\text{Sr}(\text{OH})_2$ at reflux in water for ≥ 5 h to give strontium ranelate and its hydrates. Thus, the octahydrate of strontium ranelate was prepared by this method (96% yield and 98% purity in final step).

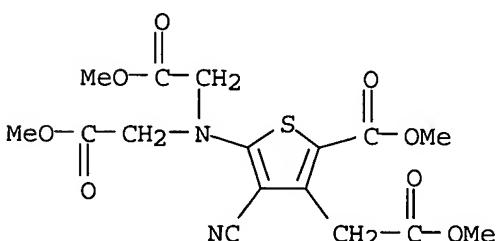
IT 674773-13-8P

RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)

(for industrial preparation of strontium ranelate and its hydrates)

RN 674773-13-8 HCPLUS

CN 3-Thiopheneacetic acid, 5-[bis(2-methoxy-2-oxoethyl)amino]-4-cyano-2-(methoxycarbonyl)-, methyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 3 OF 4 HCPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1991:492057 HCPLUS

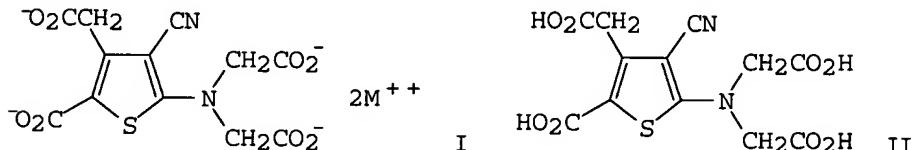
DOCUMENT NUMBER: 115:92057

TITLE: Preparation of bivalent metal salts of

[bis (carboxymethyl)amino]thiophene derivative for the
 treatment of osteoporosis and liver disease
 INVENTOR(S): Wierzbicki, Michel; Bonnet, Jacqueline; Brisset,
 Martine; Tsouderos, Yannis
 PATENT ASSIGNEE(S): ADIR et Cie., Fr.
 SOURCE: Eur. Pat. Appl., 10 pp.
 CODEN: EPXXDW
 DOCUMENT TYPE: Patent
 LANGUAGE: French
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 415850	A1	19910306	EP 1990-402401	19900831
EP 415850	B1	19940112		
R: AT, BE, CH, FR 2651497	DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE	19910308	FR 1989-11475	19890901
FR 2651497	A1	19911025		
ZA 9006716	A	19910626	ZA 1990-6716	19900823
CA 2024419	AA	19910302	CA 1990-2024419	19900831
CA 2024419	C	19990720		
AU 9062033	A1	19910307	AU 1990-62033	19900831
AU 624022	B2	19920528		
JP 03169876	A2	19910723	JP 1990-232271	19900831
JP 06092386	B4	19941116		
US 5128367	A	19920707	US 1990-576225	19900831
AT 100093	E	19940115	AT 1990-402401	19900831
ES 2062450	T3	19941216	ES 1990-402401	19900831
PRIORITY APPLN. INFO.:			FR 1989-11475	A 19890901
			EP 1990-402401	A 19900831

GI



AB The title compds. I ($M = Sr, Ca, Mg$) were prepared. Treatment of carboxylic acid II with aqueous $Sr(OH)_2$ solution gave I.4H₂O ($M = Sr$) (III). In an *in vitro*

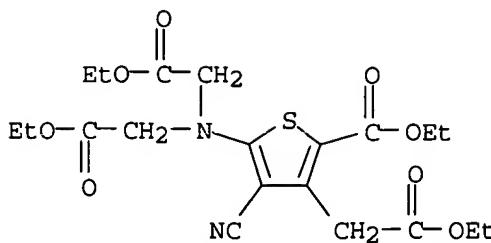
test, III at 10-4 M decreased bone resorption by about 5%.

IT 58194-26-6

RL: RCT (Reactant); RACT (Reactant or reagent)
 (saponification of)

RN 58194-26-6 HCPLUS

CN 3-Thiopheneacetic acid, 5-[bis(2-ethoxy-2-oxoethyl)amino]-4-cyano-2-(ethoxycarbonyl)-, ethyl ester (9CI) (CA INDEX NAME)



L5 ANSWER 4 OF 4 HCPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1976:59258 HCPLUS

DOCUMENT NUMBER: 84:59258

TITLE: Reactivity of 2-aminothiophenes. Application to synthesis of thieno[2,3-b]pyrroles

AUTHOR(S): Wierzbicki, Michel; Cagniant, Denise; Cagniant, Paul

CORPORATE SOURCE: Fac. Sci., Univ. Metz, Metz, Fr.

SOURCE: Bulletin de la Societe Chimique de France (1975), (7-8, Pt. 2), 1786-92

CODEN: BSCFAS; ISSN: 0037-8968

DOCUMENT TYPE: Journal

LANGUAGE: French

OTHER SOURCE(S): CASREACT 84:59258

GI For diagram(s), see printed CA Issue.

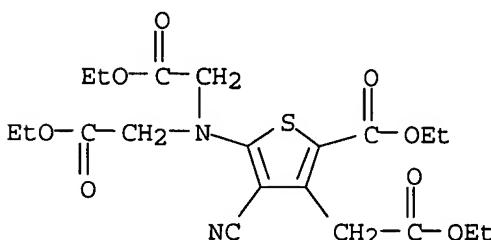
AB Thienopyrroles I (R = H, Ac; R1 = OH, NH2; R2 = CH₂CO₂Et, Me; R3 = CO₂Et, Ac) were prepared by treating the thiophenes II (R₄ = H; R₅ = CO₂Et, CN) with BrCH₂CO₂Et and Dieckmann reaction of II (R₄ = CH₂CO₂Et). I (R₁ = OH) were alkylated with BrCH₂CO₂Et or acetylated. I (R₂ = NH₂) were acetylated and diazotized.

IT 58194-26-6P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 58194-26-6 HCPLUS

CN 3-Thiopheneacetic acid, 5-[bis(2-ethoxy-2-oxoethyl)amino]-4-cyano-2-(ethoxycarbonyl)-, ethyl ester (9CI) (CA INDEX NAME)



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